## AMENDMENTS TO THE CLAIMS

1. (Currently amended) An isolated polypeptide comprising a sequence selected from the group consisting of SEQ ID No. 2 and a sequence with 95% identity thereto or SEQ ID No. 4.

- 2. (Currently amended) An isolated polynucleotide that encodes an isolated polypeptide comprising a sequence selected from the group consisting of SEQ ID No. 2 and a sequence with 95% identity thereto a polypeptide according to claim 1.
- 3. (Currently amended) An isolated polynucleotide according to claim 2, selected from the group consisting of SEQ ID No. 1, or SEQ ID No. 3, the complement of SEQ ID NO: 1, the complement of SEQ ID NO: 3, the reverse complement of SEQ ID NO: 1, the reverse complement of SEQ ID NO: 3, the reverse sequence of SEQ ID NO: 1, and the reverse sequence of SEQ ID NO: 3.
  - 4. (Canceled)
  - 5. (Currently amended) An isolated polynucleotide comprising a nucleotide sequence that differs from SEQ ID No: 1 or SEQ ID No. 3 as a result of silent substitution(s) or substitution(s) that results in conservative substitution(s) in the resulting amino acid.
  - 6. (Currently amended) An isolated polypeptide encoded by a polynucleotide of elaim 4 or claim 5 comprising a nucleotide sequence that differs from SEQ ID No: 1 or SEQ ID No. 3 as a result of silent substitution(s) or substitution(s) that results in conservative substitution(s) in the resulting amino acid.
  - 7. (Currently amended) A fusion protein comprising at least one polypeptide according to claim 1 or claim 6 or a fragment thereof and additional amino acids.
  - 8. (Currently amended) A vector comprising a polynucleotide according to any one of claims 2 or to-5.
  - 9. (Currently amended) The vector according to claim 8, further comprising, in the 5'-3' direction:
    - a) a gene promoter sequence; and
    - b) a gene termination sequence. a polynucleotide sequence according to any one of claims 2 to 5; and
      - c) a gene termination sequence.

10. (Currently amended) The vector according to claim 8-or claim-9, wherein the polynucleotide is in a sense orientation.

- 11. (Currently amended) The vector according to claim 8-or claim 9, wherein the polynucleotide is in an antisense orientation.
- 12. (Currently amended) A host cell containing a vector according to any one of claims 8-to-11.
- 13. (Currently amended) A composition for regulating muscle growth, comprising an active ingredient selected from the group consisting of any one of:
  - a) a polynucleotide comprising SEQ ID No. 1, or SEQ ID No.3, or SEQ ID NO: 5,
  - b) a fragment or variant of (a),
  - c) a polynucleotide having at least 95%, 90% or 70% sequence identity to (a),
  - d) a complement of any one of (a) to (c),
  - e) a reverse complement of any one of (a) to (c),
  - f) an antisense polynucleotide of any one of (a) to (c),
  - g) a polypeptide encoded by any one of (a) to (c),
  - h) a polypeptide comprising SEQ ID No. 2 or SEQ ID No. 4,
  - i) a fragment or variant of (g) or (h), and
  - j) a polypeptide having at least 95%, 90% or 70% sequence identity relating to (g) or (h); and

a pharmaceutically acceptable diluent, excipient or carrier.

- 14. (Canceled)
- 15. (Currently amended) A composition for modulating mighty gene expression comprising a compound capable of binding to a polynucleotide selected from the group consisting any one of:
  - a) SEQ ID No. 1, SEQ ID No. 3, or SEQ ID No. 5,
  - b) a polynucleotide that encodes a polypeptide of SEQ ID No. 2 or SEQ ID No. 4,
  - c) a polynucleotide having at least 95%, 90% or 70% sequence identity to (a) or

(b),

- d) a complement of any one of (a) to (c),
- e) a reverse complement of any one of (a) to (c), and
- f) a fragment or variant of any one of (a) to (e); and

## a pharmaceutically acceptable diluent, excipient or carrier.

- 16. (Original) The composition according to claim 15 wherein the compound is an anti-sense polynucleotide.
- 17. (Currently amended) The composition according to claim 15 or claim 16 wherein the compound is an interfering RNA molecule.
- 18. (**Original**) The composition according to claim 17 wherein the interfering RNA molecule is an RNAi or siRNA molecule.
- 19. (Original) The composition according to claim 15, wherein the compound is myostatin.
- 20. (Original) The composition according to claim 15, wherein the compound is a myostatin mimetic.
- 21. (Original) The composition according to claim 20, wherein the myostatin mimetic is a myostatin peptide C-terminally truncated at or between amino acid positions 330 and 350.
- 22. (Currently amended) The composition according to claim 20 or claim 21, wherein the myostatin mimetic is a myostatin peptide C-terminally truncated at any one a position selected from the group consisting of amino acid positions 330, 335, and 350.
- 23. (Original) The composition according to claim 15, wherein the compound is an antibody.
  - 24. (Canceled)
  - 25. (Canceled)
  - 26. (Canceled)
  - 27. (Canceled)
- 28. (Currently amended) A method of regulating muscle growth of an organism, comprising administering to said organism a composition according to any one of claims 13 or 15to 23.
- 29. (Currently amended) The method according to claim 28, for the production of an animal having increased muscle mass in said organism.
- 30. (Currently amended) The method according to claim 28, for the treatment or prophylaxis of a disease associated with muscle growth in said organism.

31. (Currently amended) The methods method according to claim 30, wherein the disease is associated with muscle atrophy.

- 32. (Currently amended) The method according to claim 30-or-claim-31, wherein the disease is selected from the group consisting of muscular dystrophy, muscle cachexia, atrophy, hypertrophy, muscle wasting associated cancer or HIV, amyotrophic lateral sclerosis (ALS), or and diseases associated with cardiac muscle growth, including infarct.
- 33. (Currently amended) A method according to claim 28, for promoting muscle regeneration after muscle injury in said organism.
  - 34. (Cancelled).
  - 35. (Cancelled).
  - 36. (Cancelled).
  - 37. (Cancelled).
  - 38. (Cancelled).
- 39. (Currently amended) A transgenic animal comprising a vector according to any one of claims 8 to 11; or a composition according to any one of claims 13 to 18.
- 40. (Original) The transgenic animal according to claim 39, wherein said animal has an increased muscle mass.
- 41. (Currently amended) The transgenic animal according to claim 39-or-claim 40, selected from the group consisting of a sheep, cow, bull, deer, poultry, turkey, pig, horse, mouse, rat or and human.
- 42. (Currently amended) A method of predicting muscle mass in an animal, comprising the steps of:

obtaining a sample from the animal,

iii) determining the gene expression level from a polynucleotide having a sequence of SEQ ID No. 1 or SEQ ID No. 3, a polynucleotide having at least 95%, 90% or 70% sequence identity to SEQ ID No. 1 or SEQ ID No. 3, or a fragment or variant thereof; or determining the amount of a polypeptide having a sequence of SEQ ID No. 2 or SEQ ID No. 4, a polypetide having at least 95%, 90% or 70% sequence identity to SEQ ID No. 2 or SEQ ID No. 4, or a fragment or variant thereof,

iv) comparing the gene expression level or amount of polypeptide to an average; and

v) predicting the muscle mass of said animal based on the gene expression level.

- 43. (Original) The method according to claim 42, wherein the level of gene expression is determined using RTPCR or northern analysis.
- 44. (Original) The method according to claim 43, wherein the amount of the polypeptide is determined using ELISA or Western blot analysis.
- 45. (Currently Amended) A method of detecting a variant of mighty, comprising the use of a nucleotide sequence selected from the group consisting of:
  - a) SEQ ID No. 1, SEQ ID No. 3, or SEQ ID No. 5,
  - b) a polynucleotide that encodes a polypeptide of SEQ ID No. 2 or SEQ ID No. 4,
  - c) a polynucleotide having at least 95%, 90% or 70% sequence identity to (a) or (b),
    - d) a complement of any one of (a) to (c),
    - e) a reverse complement of any one of (a) to (c), and
  - f) a fragment or variant of any one of (a) to (e), to screen a sample from an organism for the variant of mighty.
- 46. (Original) The method according to claim 45, wherein the variant is a polymorphism.
- 47. (Original) The method according to claim 46, wherein the polymorphism is a single nucleotide polymorphism.
- 48. (Currently amended) The method according to any one of claims 45 to 47, wherein the variant of mighty is associated with an altered muscle phenotype.
- 49. (Currently amended) A method of breeding an animal having improved muscle mass comprising the steps of:

selecting one or more animals predicted to have an increase in muscle mass using the method according to any one of claims 42 to 44 or 48, and

breeding the one or more animals predicted to have an increased muscle mass to produce an animal having an improved muscle mass.

50. (Currently amended) The method according to claim 49, wherein the animal is selected from the group consisting of a sheep, cow, bull, deer, poultry, turkey, pig, horse, mouse, rat, fish or and human.

51. (Currently Amended) An antibody that preferentially binds a polypeptide having a sequence of SEQ ID NO. 2 or SEQ ID NO. 4 or a polypeptide having at least 95%, 90% or 70% sequence identity to SEQ ID NO. 2 or SEQ ID NO. 4.

- 52. (Currently Amended) An antigenic fragment of a polypeptide comprising a sequence of SEQ ID NO.2 or SEQ ID NO. 4 in the production of an antibody that preferentially binds a sequence of SEQ ID NO. 2 or SEQ ID NO. 4 or a polypeptide having at least 95%, 90% or 70% sequence identity to SEQ ID NO. 2 or SEQ ID NO. 4.
- 53. (Currently Amended) An isolated polynucleotide selected from the group consisting comprising any one of:
  - a) a polynucleotide comprising the sequence of SEQ ID No: 5,
  - b) a polynucleotide having comprising at least 95%, 90% or 70% sequence identity to SEQ ID No. 5, and
  - c) a <u>polynucleotide comprising a fragment or variant thereof of (a) or (b)</u> having promoter activity.
  - 54. (Canceled)
- 55. (Currently amended) An isolated polynucleotide according to claim <u>53</u>54, comprising at least the 200 nucleotides upstream of the mighty initiation site.
- 56. (Currently amended) An isolated polynucleotide according to claims <u>53</u>54 or elaim <u>55</u>, comprising a fragment selected from the group consisting of those fragments of any on of 209, 287, 315, 400, 600, 1000 and 2100 nucleotides upstream of the mighty initiation site.
- 57. (Currently amended) A vector comprising a polynucleotide according to any one of claims 5354 to 56.
- 58. (Currently amended) An isolated host cell containing a vector according to claim 57.
- 59. (Currently amended) A method of screening for one or more compounds that are potentially useful in inhibiting or promoting muscle growth, comprising the steps of:

inserting a polynucleotide according to any one of claims 5354 to 56 into a suitable vector linked to a suitable marker gene;

transforming a suitable host cell with the vector; administering a compound of interest to the host cell; and determining any difference in the level of the marker gene expression.

60. (Currently amended) The method according to claim 59, wherein the vector is selected from the group consisting any one of a prokaryotic plasmid, a eukaryotic plasmid or and a viral vector.

- 61. (Currently amended) The method according to claim 59-or-elaim 60, wherein the marker gene is any one of a polynucleotide that encodes any one a protein selected from the group consisting of: a green fluorescent protein, a red fluorescent protein, a luciferase enzyme, or and a  $\beta$ -galactosidase enzyme.
- 62. (Currently amended) A method of expressing a desired protein in a muscle cell, comprising the steps of:

isolating a polynucleotide sequence that encodes the gene to be expressed; inserting a polynucleotide according to any one of claims 5354 to 56, operably linked to the polynucleotide sequence of the protein to be expressed in a 5' - 3' orientation, into a suitable vector, and

introducing the vector into a muscle host cell.

- 63. (Currently amended) The method according to claim 62, wherein the vector is any one selected from the group consisting of a eukaryotic vector, viral vector, or and any vector suitable for gene therapy.
- 64. (Currently amended) The method according to claim 62-or claim 63, wherein the host cell is any one selected from the group consisting of a primary myoblast cell line, a transformed myoblast cell line or and any cell line in which the mighty promoter is active.
- 65. (Currently amended) The method according to claim 62-or-claim 63, wherein the host cell is an *in vivo* skeletal or cardiac muscle cell of a host animal.
- 66. (Currently amended) The method according to claim 65, wherein the host animal is any one selected from the group consisting of a sheep, cow, deer, bull, poultry, turkey, pig, horse, mouse, rat, fish or and human.